

EXPRESS MAIL EM230308762US
MAY 19, 1998

APPLICATION OF QINGYUN LIU AND FANG CHEN
5 MERCK CASE NO. 19959Y

TITLE OF THE INVENTION
HUMAN UNCOUPLING PROTEIN 3

10

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a non-provisonal application of U.S.
Provisional Application Serial No.60/069,141 filed December 9, 1997,
which is a continuation-in-part of U.S. Provisional Application Serial
15 No. 60/047,179, filed May 20, 1997.

STATEMENT REGARDING FEDERALLY-SPONSORED R&D

Not applicable.

20

REFERENCE TO MICROFICHE APPENDIX

Not applicable

FIELD OF THE INVENTION

25

The present invention relates to an isolated nucleic acid
molecule (polynucleotide) which encodes a novel human uncoupling
protein, referred to throughout as uncoupling protein 3 (UCP3). The
present invention also relates to recombinant vectors and recombinant
hosts which contain a DNA fragment encoding human UCP3,
30 substantially purified forms of associated human UCP3, human UCP3
mutant proteins, and methods associated with identifying compounds
which modulate energy expenditure and body weight regulation, as well
as affecting diseases such as obesity and diabetes.

35

The balancing of energy expenditure and food intake plays a critical role in the control of body weight. Heat generation from uncoupling activity in mitochondria is responsible for a significant portion of energy expenditure, and therefore regulation of uncoupling activities provides a direct target for the treatment of obesity.

It is known that uncoupled mitochondrial respiration in brown adipose tissue (BAT) plays an important role in the regulation of energy balance in rodents.

10 Bouillaud et al. (1986, *J. Biol. Chem.* 261(4): 1487-1490) disclose a rat cDNA clone which encodes rat uncoupling protein 1 (UCP1). Rat UCP1 is a mitochondrial inner membrane protein synthesized in the cytosol from a nuclear encoded transcript which is expressed in brown adipose tissue.

15 Bouillaud et al. (1988, *Biochem. Biophys Res. Comm.* 157: 783-792) disclose a genomic clone encoding rat UCP1.

Nicholls and Locke (1984, *Physiol. Rev.* 64: 1-64) review UCP1, noting that the rat 32 kD protein forms a proton channel through the mitochondrial inner membrane and is active in uncoupling ATP synthesis from heat production in BAT.

Jacobsson et al. (1985, *J. Biol. Chem.* 260: 16250-16254) disclose cDNA clones for mRNA isolated from mouse brown adipose tissue. Mouse ucp1 mRNA was shown to be induced in brown adipose tissue by exposure to cold.

25 Kozak et al. (1988, *J. Biol. Chem.* 263: 12274-12277) disclose
the genomic clone encoding mouse UCP1.

Cassard et al. (1990, *J. Cell. Biochem.* 43: 255-264) discloses the genomic clone for human ucp1, as well as the deduced amino acid sequence for human UCP1. The authors show that rat and human UCP1 are 79% homologous at both the nucleotide and amino acid level.

UCP1 activity in thermogenesis and uncoupled energy dissipation is limited to brown adipose tissue. Therefore, it is not expected that UCP1 is actively involved in the cause and effect of body weight indications such as obesity and diabetes in vertebrates such as humans, which contain limited amount of brown fat.

5 Fleury et al. (1997, *Nature Genetics* 15: 269-272) disclose a gene encoding both a mouse and human mitochondrial uncoupling protein, designated mouse UCP2 and human UCP2, respectively. The deduced human amino acid sequence encodes a 33 kD protein which is approximately 95% homologous to the mouse protein at the amino acid level. As with human UCP1, human UCP2 comprises 3 mitochondrial carrier protein motifs and an ATP binding site. The authors show a wide range of tissue specific expression of the human ucp2 gene, including skeletal muscle, lung, heart, placenta, stomach, as well as immune systems tissue such as spleen, thymus, leukocytes, macrophages and bone marrow. The authors mapped the human ucp2 gene to chromosome 11, which has been linked to obesity and hyperinsulinaemia in the mouse.

15 It would be advantageous to identify a gene encoding an additional human uncoupling protein wherein expression is for the most part limited to skeletal muscle. A nucleic acid molecule expressing an additional human uncoupling protein in such a specific manner would be extremely useful in screening for compounds acting as a modulator of to obesity and hyperinsulinaemia. Such a compound or compounds will be useful in controlling obesity as well as deleterious indications associated with obesity, such as diabetes. Additionally, such a nucleic acid molecule will be useful in gene therapy applications to overcome the deleterious effects of obesity and obesity-related complications, such as diabetes. The present invention addresses and meets this need.

SUMMARY OF THE INVENTION

30 The present invention relates to a purified or isolated nucleic acid molecule (polynucleotide) which encodes a novel vertebrate uncoupling protein.

The present invention relates to a purified or isolated nucleic acid molecule (polynucleotide) which encodes a novel human uncoupling protein.

35 The present invention also relates to a purified or isolated nucleic acid molecule which encodes a human uncoupling

199391

protein which is predominantly expressed within human skeletal muscle.

A preferred aspect of the present invention is disclosed in Figure 1A-1B and SEQ ID NO:11, a purified human cDNA
5 encoding a novel uncoupling protein, UCP3.

Another specific embodiment of the present invention relates to isolated biologically active fragments or mutants of a nucleic acid molecule encoding human uncoupling protein 3, disclosed herein in Figure 2A-2D, Figure 3 and SEQ ID NO:12. Any
10 such polynucleotide includes but is not necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of diagnostic, therapeutic or prophylactic use.

The present invention relates to a purified or isolated nucleic acid molecule (polynucleotide) which encodes a novel mouse uncoupling protein, as set forth as SEQ ID NO:17.
15

The present invention also relates to a purified or isolated nucleic acid molecule which encodes a mouse uncoupling protein which is predominantly expressed within human skeletal muscle, the expressed protein as set forth in SEQ ID NO:18.
20

The purified nucleic acid molecule of the present invention may include a deoxyribonucleic acid molecule (DNA), such as genomic DNA and complementary DNA (cDNA), which may be
25 single (coding or noncoding strand) or double stranded, as well as synthetic DNA, such as a synthesized, single stranded polynucleotide. The isolated nucleic acid molecule of the present invention may also include a ribonucleic acid molecule (RNA).

The present invention also relates to recombinant
30 vectors and recombinant hosts, both prokaryotic and eukaryotic, which contain the substantially purified mammalian nucleic acid molecules disclosed throughout this specification.

The present invention also relates to subcellular membrane fractions of the recombinant host cells (both prokaryotic
35 and eukaryotic as well as both stably and transiently transformed

5 The present invention also relates to a substantially purified human uncoupling protein wherein the native form is substantially localized within mitochondria of skeletal muscle.

Another preferred aspect of the present invention is disclosed in Figure 7 and SEQ ID NO:18, the amino acid sequence of the exemplified mouse uncoupling protein, mUCP3.

20 The present invention also relates to methods of expressing the UCP proteins or protein fragments disclosed herein, assays employing these human UCPs, cells expressing these UCPs, and compounds identified through the use of these UCPs, including modulators of energy expenditure and body weight regulation, either through direct
25 contact with the mitochondrial uncoupling protein, or a compound which acts in trans to modulate UCP expression. Such modulators identified in this process are useful as therapeutic agents for controlling obesity, diabetes and other related diseases.

Figure 1A and Figure 1B show the nucleotide sequence (SEQ ID NO:11) which comprises the full-length cDNA encoding human uncoupling protein 3.

Figure 2A, Figure 2B, Figure 2C and Figure 2D show the translation of the open reading frame (SEQ ID NO:12) of the full-length human uncoupling protein 3 nucleotide sequence.

Figure 3 shows the amino acid sequence of human uncoupling protein 3 (SEQ ID NO:12).

Figure 4A, Figure 4B and Figure 4C show Northern blot analysis of human uncoupling protein 3 in human tissue.

Figure 5 shows the orientation of vector and gene specific primers used to screen a human fetal brain cDNA library for a full-length human UCP3 cDNA.

Figure 6A and 6B shows the nucleotide sequence (SEQ ID NO:17) of a cDNA which encodes the full-length mouse uncoupling protein 3.

Figure 7 shows the amino acid sequence (SEQ ID NO:18) of the mouse uncoupling protein 3.

Figure 8A, Figure 8B and Figure 8C show flow cytometry analysis of mitochondria membrane potential with UCP3 expression in yeast. *TheS. cerevisiae* strain W303 containing vector alone (Figure 8A) or UCP3 expression plasmids (Figure 8B and Figure 8C) were stained with the potential sensitive dye, DiOC6 and analyzed by FACS. The X-axis represents the intensity of fluorescence on logarithmic scale while the Y-axis indicates the number of cells. "+gal" indicates that cells were induced with galactose for 5 hours. "-gal" indicates cells were maintained in raffinose media without the induction by galactose. A decrease in fluorescence intensity indicates a reduction in mitochondria membrane potential.

Figure 9A and Figure 9B shows the response of UCP3 expression to adenovirus-mediated leptin treatment. Figure 9A shows measurement of weight loss. Y-axis indicates the average weight expressed in percentage of weight relative to the day of injection (Day 0). Figure 9B shows UCP3 mRNA level. Y-axis indicates the relative level of UCP3 mRNA (the average of the Adb-gal and control animals is assigned a value of 1). Standard deviation for each group is as marked. Ad-Leptin: first generation E1-deleted, replication deficient adenoviral vector expressing leptin; HD-leptin: helper-dependent adenoviral vector

expressing leptin; Adb-gal: first generation adenoviral vector expressing b-galactosidase; Control: untreated.

DETAILED DESCRIPTION OF THE INVENTION

5 The present invention relates to isolated nucleic acid and protein forms which represent a vertebrate uncoupling protein, preferably human and mouse forms. The isolated genes and concomitant translation products which make up a portion of the present invention may be used in various ways in the treatment of
10 obesity and other related diseases, including but not limited to diabetes. The uncoupling protein of the present invention is a protein which is translated in the cytosol from a nuclear encoded mRNA transcript. This uncoupling protein is presumably imported into the mitochondrial inner membrane post-translationally.

15 The process of oxidative phosphorylation pumps protons to the outside of the inner membrane of the mitochondria, leading to the generation of a membrane potential across the inner membrane. This membrane potential is then used to drive ATP synthesis. The purified uncoupling proteins of the present invention have the ability to uncouple
20 the protonmotive force associated with oxidative phosphorylation, resulting in the concomitant release of large amounts of energy. It will be within the scope of the present invention to use the herein disclosed nucleic acids to construct recombinant vectors and recombinant host cells wherein the human UCP3 gene is expressed, either stably or
25 transiently. Such transformed recombinant cell lines will be useful in screening for modulators of UCP3 activity, and hence, as effectors of energy expenditure and body weight regulation. The isolated nucleic acid fragments of the present invention will also be useful in gene therapy applications regarding therapeutic treatment of obesity and
30 obesity related indications, including but not limited to diabetes, as well as conditions such as mitochondrial-associated hypermetabolism.

To this end, the present invention relates to a purified or isolated nucleic acid molecule (polynucleotide) which encodes a novel human uncoupling protein.

The present invention also relates to a purified or isolated nucleic acid molecule which encodes a human uncoupling protein which is predominantly expressed within human skeletal muscle.

5 A preferred aspect of the present invention is disclosed in Figure 1 and SEQ ID NO:11, a purified human cDNA encoding a novel uncoupling protein, UCP3, which is as follows:

TCGAACTCAC TCACCTCCCC TCTCACCTCA CTGCCCTCAC CAGCCAGCCT
CTTGTCAAGT GATCAGGCTG TCAACCAACT TCTCTAGGAT AAGGTTTCAG
10 GTCAGCCTGT GTGTATAAGA CCAGTGCCAA GCCAGAAGCA GCAGAGACAA
CAGTGAATGA CAAGGAGGGG CCATCCAATC CCTGCTGCCA CCTCCTGGGA
TGGAGCCCTA GGGAGCCCCCT GTGCTGCCCC TGCCGTGGCA GGA CTACAG
CCCCACCGCT GCACTGAAGC CCAGGGCTGT GGAGCAGCTC TCTCCTTGGA
CTCCTCTCGG CCCTAAAGGG ACTGGGCAGA GCCTTCCAGG ACTATGGTTG
15 GACTGAAGCC TTCAGACGTG CCTCCCACCA TGGCTGTGAA GTTCCTGGGG
GCAGGCACAG CAGCCTGTTT TGCTGACCTC GTTACCTTTC CACTGGACAC
AGCCAAGGTC CGCCTGCAGA TCCAGGGGGA GAACCAGGCG GTCCAGACGG
CCCGGCTCGT GCAGTACCGT GGCGTGCTGG GCACCATCCT GACCATGGTG
CGGACTGAGG GTCCCTGCAG CCCCTACAAT GGGCTGGTGG CCGGCCTGCA
20 GCGCCAGATG AGCTTCGCCT CCATCCGCAT CGGCCTTTAC GACTCCGTCA
AGCAGGTGTA CACCCCCAAA GGC GCGGACA ACTCCAGCCT CACTACCCGG
ATTTTGGCCG GCTGCACCAC AGGAGCCATG GCGGTGACCT GTGCCCAGCC
CACAGATGTG GTGAAGGTCC GATTTTCAGGC CAGCATACAC CTCGGGCCAT
CCAGGAGCGA CAGAAAATAC AGCGGGACTA TGGACGCCTA CAGAACCATC
25 GCCAGGGAGG AAGGAGTCAG GGGCCTGTGG AAAGGAACTT TGCCCAACAT
CATGAGGAAT GCTATCGTCA ACTGTGCTGA GGTGGTGACC TACGACATCC
TCAAGGAGAA GCTGCTGGAC TATCACCTGC TCACTGACAA CTTCCCCCTGC
CACTTTGTCT CTGCCTTTGG AGCCGGCTTC TGTGCCACAG TGGTGGCCTC
CCCGGTGGAC GTGGTGAAGA CCCGGTATAT GAACTCACCT CCAGGCCAGT
30 ACTTCAGCCC CCTCGACTGT ATGATAAAGA TGGTGGCCCA GGAGGGCCCC
ACAGCCTTCT ACAAGGGATT TACACCCTCC TTTTGTGCGTT TGGGATCCTG
GAACGTGGTG ATGTTTCGTAA CCTATGAGCA GCTGAAACGG GCCCTGATGA
AAGTCCAGAT GTTACGGGAA TCACCGTTTT GAACAAGACA AGAAGGCCAC
TGGTAGCTAA CGTGTCCGAA ACCAGTTAAG AATGGAAGAA AACGGTGCAT
35 CCACGCACAC ATGGACACAG ACCCACACAT GTTTACAGAA CTGTTGTTTA

CTTGTTGCTG ATTCAAGAAA CAGAAGTAGA AGAGAGAGGA TTCTGGTCTT
 CACTGCCATG CCTCAAGAAC ACCTTTGT TT TGCAC TGACA AGATGGAAAA
 TAAATTATAT TAATTTTGA AACCCATTAG GCATGCCTAA TATTTAGGCA
 AGAGAAAATA AACCAAGATA GATCCATTG GACAAAATGG AAGGTGGAG
 5 ACGTGTATCC CCGTGAAATC TGGTCAGATA ATGAATGATA AGCAGGAAGG
 ATGGCAAGCA CGGGACAGGA GGGGCCACA ATGGAGTGGG AGATCAGCCA
 CGGAGCCTGG AGGGACGCCA CCCAGCAACA CAGAGCTGGC GACTGCAGCT
 GCACCATCAC ACATGCATCA TCAGCCTATT TGTAATATGT CTGCCACAGA
 GAGTCCTTTG GGATTCTAGG AAACCCAAGG AACAGAGAA AAAACTAGAG
 10 CCTGTGCTAA AGAAGCCTGC TGGGCCCATG TGAGGCTGGG GTCGTAAATA
 TTCCCCGACG ACACTGAAGA ATCAAGAGGG CAGCCCCCAC TTCTCCTACA
 AAATGGAGGG AGCCATCCCT TCCCTGTCCA CCTCACCAGG GGTGCTATGA
 CATGCAAGTG AGAAGCTGGG CATGAACGCA CTTTATAAAA GCAAAAGCTC
 TGTGTAAATC TAACTACAAG GACAATGCCT TGGGAGAGAT TTTGTCGGGA
 15 CAGAGAGGAG TTGCCAGGGA AGAAGGTTTG AAAGATACGG TTGTCTAGAG
 GTGAGACCAA AGGATCCAGA GACTTGGGGA CCAGAGGTGA CAGTGGATGA
 CGTGAAGCCA CAGGAGCCCC ACCCCCATGC AGCTTTTTC CCACCCCCC
 CACCACGCGC TCAATCATGA GTACCTCAA GGATTGTTGG GCTTGGGGGA
 AAAGAGGTGG ATTCCTGGGC AAGAACCTAA AGTAGCAGGA (SEQ ID NO:11).

20 Another specific embodiment of the present invention
 relates to isolated biologically active fragments or mutants of a
 nucleic acid molecule encoding human uncoupling protein 3,
 disclosed herein as SEQ ID NO:11 (see also Figure 1A-1B). Any such
 polynucleotide includes but is not necessarily limited to nucleotide
 25 substitutions, deletions, additions, amino-terminal truncations and
 carboxy-terminal truncations such that these mutations encode
 mRNA which express a protein or protein fragment of diagnostic,
 therapeutic or prophylactic use.

The purified nucleic acid molecule of the present
 30 invention may include a deoxyribonucleic acid molecule (DNA), such
 as genomic DNA and complementary DNA (cDNA), which may be
 single (coding or noncoding strand) or double stranded, as well as
 synthetic DNA, such as a synthesized, single stranded
 polynucleotide. The isolated nucleic acid molecule of the present
 35 invention may also include a ribonucleic acid molecule (RNA).

The present invention also relates to recombinant vectors and recombinant hosts, both prokaryotic and eukaryotic, which contain the substantially purified nucleic acid molecules disclosed throughout this specification.

5 The present invention also relates to a substantially purified human uncoupling protein wherein the native form is substantially localized within mitochondria of skeletal muscle.

A preferred aspect of the present invention is disclosed in Figure 2A-2D, Figure 3 and SEQ ID NO:12, the amino acid
10 sequence of the exemplified human uncoupling protein, UCP3, which is as follows:

MVGLKPSDVPPTMAVKFLGAGTAACFADLVTFPLDTAKVRLQIQGENQAVQTARLV
QYRGVLGTILTMVRTEGPCSPYNGLVAGLQRQMSFASIRIGLYDSVKQVYTPKGAD
NSSLTTRILAGCTTGAMAVTCAQPTDVVKVRFQASIHGSPSRSDRKYSGMTDAYRT
15 IAREEGVRGLWKGTLPNIMRNAIVNCAEVVITYDILKEKLLDYHLLTDNFPCHFVSA
FGAGFCATVVASPVDVVKTRYMNSPPGQYFSPDCMIKMVAQEGPTAFYKGFTPSF
LRLGSWNVVMFVITYEQLKRALMKVQMLRESPF, as set forth in three-letter
abbreviation in SEQ ID NO:12.

The present invention also relates to biologically active
20 fragments and/or mutants of a novel human uncoupling protein, including but not necessarily limited to amino acid substitutions, deletions, additions, amino terminal truncations and carboxy-terminal truncations such that these mutations provide for proteins or protein fragments of diagnostic, therapeutic or prophylactic use.

25 The present invention also relates to methods of expressing the vertebrate UCP proteins or protein fragments disclosed herein, assays employing these human and mouse UCPs, cells expressing these UCPs, and compounds identified through the use of these UCPs, including modulators of energy expenditure and body weight regulation, either
30 through direct contact with the mitochondrial uncoupling protein, or a compound which acts in *trans* to modulate ucp expression. Such modulators identified in this process are useful as therapeutic agents for controlling obesity, diabetes and other related diseases.

Other uses of the nucleic acid fragment and protein fragments of
35 the present invention include, but are not limited to, identifying trans-

Therefore, the present invention also relates to methods of expressing a nucleic acid fragment encoding UCP, including but not limited to the exemplified human and/or mouse *ucp3* or gene fragments disclosed herein, recombinant prokaryotic and eukaryotic expression vectors containing such a nucleic acid fragment, recombinant host cells transfected with these vectors wherein a *ucp* gene or gene fragment is expressed at predetermined levels through appropriate vector construction, and compounds identified through the interaction with either regulatory regions of a *ucp* gene or a compound which interacts with the mature, mitochondrial-localized form of a UCP protein. Such a compound may be useful in treating disorders associated with regulation of body weight and concomitant intake and expenditure of energy (or lack thereof), including but not limited to obesity and diabetes.

- 11 -

12

T,0130

566F59 264T9060

5 CCAGGAACAG CAGAGACAAC AGTGAATGGT GAGGCCCCGGC CGTCAGATCC
 TGCTGCTACC TAATGGAGTG GATCCTTAGG GTGGCCCTGC ACTACCCAAC
 CTTGGCTAGA CGCACAGCTT CCTCCCTGAA CTGAAGCAAA AGATTGCCAG
 GCAAGCTCTC TCCTCGGACC TCCATAGGCA GCAAAGGAAC CAGGCCCCATT
 CCCCCGGGACC ATGGTTGGAC TTCAGCCCTC CGAAGTGCCT CCCACAACGG
 TTGTGAAGTT CCTGGGGGCC GGCAC TGCGG CCTGTTTTGC GGACCTCCTC
 ACTTTTCCCC TGGACACCGC CAAGGTCCGT CTGCAGATCC AAGGGGAGAA
 CCCAGGGGCT CAGAGCGTGC AGTACCGCGG TGTGCTGGGT ACCATCCTGA
 CTATGGTGCG CACAGAGGGT CCCCCGAGCC CCTACAGCGG ACTGGTCGCT

10 GGCTGCACC GCCAGATGAG TTTTGCCTCC ATTCGAATTG GCCTCTACGA
 CTCTGTCAAG CAGTTCTACA CCCCCAAGGG AGCGGACCAC TCCAGCGTCG
 CCATCAGGAT TCTGGCAGGC TGCACGACAG GAGCCATGGC AGTGACCTGC
 GCCCAGCCCA CGGATGTGGT GAAGGTCCGA TTTCAAGCCA TGATACGCCT
 GGGAAGTGA GGAGAGAGGA AATACAGAGG GACTATGGAT GCCTACAGAA

15 CCATCGCCAG GGAGGAAGGA GTCAGGGGCC TGTGGAAAGG GACTTGGCCC
 AACATCACAA GAAATGCCAT TGTCAACTGT GCTGAGATGG TGACCTACGA
 CATCATCAAG GAGAAGTTGC TGGAGTCTCA CCTGTTTACT GACAACTTCC
 CCTGTCACTT TGTCTCTGCC TTTGGAGCTG GCTTCTGTGC CACAGTGGTG
 GCCTCCCCGG TGGATGTGGT AAAGACCCGA TACATGAACG CTCCCCTAGG

20 CAGGTACCGC AGCCCTCTGC ACTGTATGCT GAAGATGGTG GCTCAGGAGG
 GACCCACGGC CTTCTACAAA GGATTTGTGC CCTCCTTTCT GCGTCTGGGA
 GCTTGGAAACG TGATGATGTT TGTAACATAT GAGCAACTGA AGAGGGCCTT
 AATGAAAGTC CAGGTACTGC GGAATCTCC GTTTTGAACA AGGCAAGCAG
 GCTGCCTGGA ACAGAACAAA GCGTCTCTGC CCTGGGGACA CAGGCCCACA

25 CGGTCCAGAA CCCTGCACTG CTGCTGACAC GAGAACTGA ACTAAAAGAG
 GAGAGTTTTA GTCCCTCCGTG TTTTCGTCTTA AAACACCTCT GTTTTGCCT
 GACCTGATGG GAAATAAATT ATATTAATTT TTAAACCCTT TCCGGTTGGA
 TGCCTAACAT TTAGGCAAGA GACAACAAAG AAAACCAGAG TCAACTCCCT
 TGAAATGTAG GAATAAAGGA TGCATAATAA ACAGGAAAGG CACAGGTTTT

30 GAGAAGATCA GCCCACAGTG TTGTCCTTGA ATCAAACAAA ATGGTCGGAG
 GAACCCTTCG GGTTCAGCAC AAAGAGGTGA CTACAGCCTT TTGGTCACCA
 GATGACTCCG CCCCTTTGTA ATGAGTCTGC CAAGTAGACT CTATCAAGAT
 TCTGGGGAAA GGAGAAAGAA CACATTGACC TGCCCCGGCG GCCGCTCGAG
 CCCTATGA, disclosed as SEQ ID NO:17.

199591

The present invention also relates to mouse forms of isolated biologically active fragments or mutants of a nucleic acid molecule encoding human uncoupling protein 3, disclosed herein as SEQ ID NO:17 (see also Figure 6A-6B). Any such polynucleotide
5 includes but is not necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of diagnostic, therapeutic or prophylactic use.

10 The purified mouse nucleic acid molecules of the present invention may include a deoxyribonucleic acid molecule (DNA), such as genomic DNA and complementary DNA (cDNA), which may be single (coding or noncoding strand) or double
15 stranded, as well as synthetic DNA, such as a synthesized, single stranded polynucleotide. The isolated nucleic acid molecule of the present invention may also include a ribonucleic acid molecule (RNA).

20 The present invention also relates to recombinant vectors and recombinant hosts, both prokaryotic and eukaryotic, which contain the substantially purified mouse nucleic acid molecules disclosed throughout this specification.

The present invention also relates to a substantially purified mouse uncoupling protein wherein the native form is substantially localized within mitochondria of skeletal muscle.

25 A preferred aspect of the present invention is disclosed in Figure 7 and SEQ ID NO:18, the amino acid sequence of the exemplified mouse uncoupling protein, UCP3, which is as follows:
MVGLQPSEVP PTTVVKFLGA GTAACFADLL TFPLDTAKVR LQIQGENPGA
QSVQYRGVLG TILTMVRTEG PRSPYSGLVA GLHRQMSFAS IRIGLYDSVK
30 QFYTPKGADH SSVAIRILAG CTTGAMAVTC AQPTDVVKVR FQAMIRLG TG
GERKYRG TMD AYRTIAREEG VRGLWKG TWP NITRNAIVNC AEMVTYDIK
EKLLESHLFT DNFPCHFVSA FGAGFCATVV ASPVDVVKTR YMNAPLGRYR
SPLHCMLKMV AQEGPTAFYK GFVPSFLRLG AWNVMMFVTY EQLKRALMKV
QVLRESPF*, as set forth in three-letter abbreviation in SEQ ID NO:18.

133031

The present invention also relates to biologically active fragments and/or mutants of a novel mouse uncoupling protein, including but not necessarily limited to amino acid substitutions, deletions, additions, amino terminal truncations and carboxy-terminal truncations such that these mutations provide for proteins or protein fragments of diagnostic, therapeutic or prophylactic use.

It is known that there is a substantial amount of redundancy in the various codons which code for specific amino acids. Therefore, this invention is also directed to those DNA sequences which contain alternative codons which code for the eventual translation of the identical amino acid. For purposes of this specification, a sequence bearing one or more replaced codons will be defined as a degenerate variation. Also included within the scope of this invention are mutations either in the DNA sequence or the translated protein which do not substantially alter the ultimate physical properties of the expressed protein. For example, substitution of valine for leucine, arginine for lysine, or asparagine for glutamine may not cause a change in functionality of the polypeptide.

It is known that DNA sequences coding for a peptide may be altered so as to code for a peptide having properties that are different than those of the naturally-occurring peptide. Methods of altering the DNA sequences include, but are not limited to site directed mutagenesis. Examples of altered properties include but are not limited to changes in the affinity of an enzyme for a substrate or a receptor for a ligand.

As used herein, "purified" and "isolated" are utilized interchangeably to stand for the proposition that the nucleic acid, protein, or respective fragment thereof in question has been substantially removed from its *in vivo* environment so that it may be manipulated by the skilled artisan, such as but not limited to nucleotide sequencing, restriction digestion, site-directed mutagenesis, and subcloning into expression vectors for a nucleic acid fragment as well as obtaining the protein or protein fragment in pure quantities so as to afford the opportunity to generate polyclonal antibodies, monoclonal antibodies, amino acid sequencing, and peptide digestion. Therefore,

the nucleic acids claimed herein may be present in whole cells or in cell lysates or in a partially purified or substantially purified form. A nucleic acid is considered substantially purified when it is purified away from environmental contaminants. Thus, a nucleic acid sequence
5 isolated from cells is considered to be substantially purified when purified from cellular components by standard methods while a chemically synthesized nucleic acid sequence is considered to be substantially purified when purified from its chemical precursors.

As used herein, "UCP3" may refer to any vertebrate form of UCP3,
10 including but not limited to mouse or human UCP3.

As used herein, "BAT" means brown adipose tissue.

As used herein, "EST" means expressed sequence tag.

As used herein, a "biologically active equivalent" or "functional derivative" of a wild type UCP3 possesses a biological
15 activity that is substantially similar to the biological activity of the wild type UCP3. The term "functional derivative" is intended to include the "fragments," "mutants," "variants," "degenerate variants," "analogs" and "homologues" or to "chemical derivatives" of the wild type UCP3 protein. The term "fragment" is meant to refer to
20 any polypeptide subset of wild type UCP3. The term "mutant" is meant to refer to a molecule that may be substantially similar to the wild type form but possesses distinguishing biological characteristics. Such altered characteristics include but are in no way limited to altered substrate binding, altered substrate affinity
25 and altered sensitivity to chemical compounds affecting biological activity of the UCP3 or UCP3 functional derivative. The term "variant" is meant to refer to a molecule substantially similar in structure and function to either the entire wild type protein or to a fragment thereof. A molecule is "substantially similar" to a wild type
30 UCP3-like protein if both molecules have substantially similar structures or if both molecules possess similar biological activity. Therefore, if the two molecules possess substantially similar activity, they are considered to be variants even if the structure of one of the molecules is not found in the other or even if the two amino acid
35 sequences are not identical.

The term "analog" refers to a molecule substantially similar in function to either the entire wild type UCP3-like protein or to a fragment thereof.

- Any of a variety of procedures may be used to clone
- 5 UCP3. These methods include, but are not limited to, (1) a RACE PCR cloning technique (Frohman, et al., 1988, *Proc. Natl. Acad. Sci.* 85: 8998-9002). 5' and/or 3' RACE may be performed to generate a full length cDNA sequence. This strategy involves using gene-specific oligonucleotide primers for PCR amplification
- 10 of UCP3 cDNA. These gene-specific primers are designed through identification of an expressed sequence tag (EST) nucleotide sequence which has been identified by searching any number of publicly available nucleic acid and protein databases; (2) direct functional expression of the UCP3 cDNA following the
- 15 construction of an UCP3-containing cDNA library in an appropriate expression vector system; (3) screening a UCP3-containing cDNA library constructed in a bacteriophage or plasmid shuttle vector with a labeled degenerate oligonucleotide probe designed from the amino acid sequence of the UCP3 protein;
- 20 (4) screening a UCP3-containing cDNA library constructed in a bacteriophage or plasmid shuttle vector with a partial cDNA encoding the UCP3 protein. This partial cDNA is obtained by the specific PCR amplification of UCP3 DNA fragments through the design of degenerate oligonucleotide primers from the amino acid
- 25 sequence known for other UCP3 kinases which are related to the UCP3 protein; (5) screening an UCP3-containing cDNA library constructed in a bacteriophage or plasmid shuttle vector with a partial cDNA encoding the UCP3 protein. This strategy may also involve using gene-specific oligonucleotide primers for PCR
- 30 amplification of UCP3 cDNA identified as an EST as described above; or (6) designing 5' and 3' gene specific oligonucleotides using SEQ ID NOS 1 or 2 as a template so that either the full length cDNA may be generated by known RACE techniques, or a portion of the coding region may be generated by these same

known RACE techniques to generate and isolate a portion of the coding region to use as a probe to screen one of numerous types of cDNA and/or genomic libraries in order to isolate a full length version of the nucleotide sequence encoding UCP3.

5 It is readily apparent to those skilled in the art that other types of libraries, as well as libraries constructed from other cells types or species types, may be useful for isolating a UCP3-encoding DNA or a UCP3 homologue. Other types of libraries include, but are not limited to, cDNA libraries derived from other
10 cells or cell lines other than human cells or tissue such as murine cells, rodent cells or any other such vertebrate host which may contain a UCP3-encoding DNA. Additionally a UCP3 gene may be isolated by oligonucleotide- or polynucleotide- based hybridization screening of a vertebrate genomic library, including
15 but not limited to a human genomic library, a murine genomic library and a rodent genomic library, as well as concomitant human genomic DNA libraries.

 It is readily apparent to those skilled in the art that suitable cDNA libraries may be prepared from cells or cell lines
20 which have UCP3 activity. The selection of cells or cell lines for use in preparing a cDNA library to isolate a UCP3 cDNA may be done by first measuring cell associated UCP3 activity using any known assay for UCP3 activity.

 Preparation of cDNA libraries can be performed by
25 standard techniques well known in the art. Well known cDNA library construction techniques can be found for example, in Sambrook, et al., 1989, *Molecular Cloning: A Laboratory Manual*; Cold Spring Harbor Laboratory, Cold Spring Harbor, New York. Complementary DNA libraries may also be obtained from
30 numerous commercial sources, including but not limited to Clontech Laboratories, Inc. and Stratagent.

 It is also readily apparent to those skilled in the art that DNA encoding UCP3 may also be isolated from a suitable genomic DNA library. Construction of genomic DNA libraries
35 can be performed by standard techniques well known in the art.

Well known genomic DNA library construction techniques can be found in Sambrook, et al., *supra*.

In order to clone the UCP3 gene by one of the preferred methods, the amino acid sequence or DNA sequence of UCP3 or a homologous protein may be necessary. To accomplish this, the UCP3 or a homologous protein may be purified and partial amino acid sequence determined by automated sequenators. It is not necessary to determine the entire amino acid sequence, but the linear sequence of two regions of 6 to 8 amino acids can be determined for the PCR amplification of a partial UCP3 DNA fragment. Once suitable amino acid sequences have been identified, the DNA sequences capable of encoding them are synthesized. Because the genetic code is degenerate, more than one codon may be used to encode a particular amino acid, and therefore, the amino acid sequence can be encoded by any of a set of similar DNA oligonucleotides. Only one member of the set will be identical to the UCP3 sequence but others in the set will be capable of hybridizing to UCP3 DNA even in the presence of DNA oligonucleotides with mismatches. The mismatched DNA oligonucleotides may still sufficiently hybridize to the UCP3 DNA to permit identification and isolation of UCP3 encoding DNA. Alternatively, the nucleotide sequence of a region of an expressed sequence may be identified by searching one or more available genomic databases. Gene-specific primers may be used to perform PCR amplification of a cDNA of interest from either a cDNA library or a population of cDNAs. As noted above, the appropriate nucleotide sequence for use in a PCR-based method may be obtained from SEQ ID NOS: 1 or 2, either for the purpose of isolating overlapping 5' and 3' RACE products for generation of a full-length sequence coding for UCP3, or to isolate a portion of the nucleotide sequence coding for UCP3 for use as a probe to screen one or more cDNA- or genomic-based libraries to isolate a full-length sequence encoding UCP3 or UCP3-like proteins.

1993A

5
10
15
20

In an exemplified method, the human UCP3 full length cDNA of the present invention was generated by a novel method of cDNA screening. Briefly, the extension of partial cDNA sequences have historically been achieved with one or both of the two commonly used methods: filter screening of cDNA libraries by hybridization with labeled probes, and 5'- and 3'-RACE with total cellular mRNA by PCR. The first method is effective but laborious and slow while the latter method is fast but limited in efficiency. This RACE protocol is hindered by limited length of extension due to the use of the entire cellular mRNA population in a single reaction. Since smaller fragments are amplified much more efficiently than larger fragments by PCR in the same reaction, PCR products obtained using the second method are often quite small. This portion of the present invention marks an improvement upon known methods of cDNA library screening by initially constructing and subdividing cDNA libraries followed by isolating 5'- and 3'- flanking fragments by PCR. Since each pool is unlikely to contain more than one clone for a given gene which is low to moderately expressed, competition between large and small PCR products in one pool does not exist, making it possible to isolate fragments of various sizes. One definite advantage of the method disclosed in this specification is the efficiency, throughput, and its potential to isolate alternatively spliced cDNA forms.

25
30
35

Therefore, this portion of the invention relates to a process for rapid extension of a partial cDNA sequence based on subdividing a primary cDNA library and DNA amplification by polymerase chain reaction (PCR). A cDNA library is constructed with cDNA primed by random, oligo-dT or a combination of both random and oligo-dT primers and then subdivided into pools at approximately 10,000 -20,000 clones per pool. Each pool is amplified separately and therefore represents an independent portion of the cDNA molecules from the original mRNA source. Samples from all the pools are collected and transferred into 96-well plates. To extend a partial cDNA sequence, such as SEQ ID NO:1 or 2, positive pools containing the partial cDNA sequence are first identified by PCR with a pair of primers complementary to the cDNA sequence. Each positive pool in the library contains an independent

clone of the cDNA sequence; within each clone are embedded the partial cDNA sequence and its flanking fragments. The flanking fragments are isolated by PCR with primers complementary to the known vector and cDNA sequences and then sequenced directly. The DNA sequences from these fragments plus the original partial cDNA sequence are assembled into a continuous fragment, resulting in the extension of the partial cDNA sequence and the eventual determination of its full-length gene sequence by repeating the process, if necessary, until a complete open reading frame is obtained.

10 The fundamental principle of this process is to subdivide a complex library into pools of about 10,000 to about 20,000 clones. A library of two million primary clones, a number large enough to cover most mRNA transcripts expressed in the tissue, can be subdivided into 188 pools and stored in two 96-well plates. Since the number of transcripts for most genes is fewer than one copy per ~10,000 transcripts in total cellular mRNA, each pool is unlikely to contain more than one clone for a given cDNA sequence. Such reduced complexity makes it possible to use PCR to isolate flanking fragments of partial cDNA sequences larger than those obtained by known methods.

20 The foregoing process is exemplified for isolation and characterization of a full-length cDNA encoding human UCP3. Briefly, a random and oligo dT primed fetal brain cDNA library consisting of approximately 4 million primary clones was constructed in the plasmid vector pBluescript (Stratagene, LaJolla, CA). The primary clones were subdivided into 188 pools with each pool containing ~20,000 clones. Each pool was amplified separately and the resulting plasmid pools were collected and transferred into two 96-well plates. Primer pairs from the 5' and 3' portion of SEQ ID NO:1 (EST-AA192136), 5' and 3' oligonucleotides from SEQ ID NO:4 (EST-AA192553) as well as oligonucleotide primers both 5' and 3' of the polylinker sequence of the vector (in this case, pBluescript SK⁻) were used to scan this human fetal brain cDNA library distributed in a 96-well plate separately. Initial positive pools were identified with the 5' and 3' primers from AA192136 (SEQ ID NOS: 5 and 6) and the pBluescript primers (SEQ ID NOS:9 and 10). The positive pools were scanned again by PCR with both pBluescript

21

0508177 051998

primers as well as the 5' and 3' primers associated with the 3' EST, AA192553 (primers 3U3F and 3U3R). This step showed that scanning with 838F (pBSK⁻, SEQ ID NO: 9) and 3U3F (hybridizing to the 3' EST AA192553; SEQ ID NO:8), generated a 2.4 Kb cDNA fragment. This
5 cDNA fragment was subcloned into the vector pTA2.1 (Invitrogen, San Diego, CA, USA) by TA cloning. Four positive clones, the original PCR fragment, as well the I.M.A.G.E. clone #628529 (SEQ ID NOS:1 and 2) were sequenced by primer walking. The sequences were then assembled into a contig of 2340 base pairs which is as set forth in SEQ ID
10 NO:11 and Figure 1A and Figure 1B. This sequence contains an open reading frame that encodes a polypeptide of 312 amino acids as set forth in SEQ ID NO:12 and Figures 2A-2D and Figure 3.

The cDNA as set forth in SEQ ID NO:11 was identified as a full-length cDNA fragment from a single positive pool. Figure 5 shows the
15 vector and EST (gene) specific primers used to isolate the human UCP3 gene. In this instance, the entire coding region and 3' untranslated region was retrieved by the 838F and 3U3F primer combination. However, the skilled artisan, aided with this specification, will understand the more far reaching cDNA cloning process disclosed
20 herein: multiple primer combinations from an EST or other partial cDNA sequence, in combination with flanking vector primer oligonucleotides may be used to "walk" in both directions away from the internal gene specific sequence and respective primers such that a contig representing a full length cDNA may be constructed. This
25 procedure relies on the ability to screen multiple pools which comprise a representative portion of the total cDNA library. This procedure is not dependent upon using a cDNA library with directionally cloned inserts. Instead, both 5' and 3' vector and gene specific primers are added and a contig map is constructed from additional screening of positive pools
30 using both vector primers and gene specific primers. Of course, these gene specific primers are initially constructed from a known nucleic acid fragment such as an expressed sequence tag. However, as the walk continues, gene specific primers are utilized from the 5' and 3' boundaries of the newly identified regions of the cDNA. As the walk

continues, there is still no requirement that the vector orientation of a yet unidentified fragment be known. Instead, all combinations are tested on a positive pool and the actual vector orientation is determined by the ability of certain vector/gene specific primers to generate the predicted PCR fragment. A full-length cDNA may then be easily constructed by known subcloning procedures.

A variety of mammalian expression vectors may be used to express recombinant UCP3 in mammalian cells. Expression vectors are defined herein as DNA sequences that are required for the transcription of cloned DNA and the translation of their mRNAs in an appropriate host. Such vectors can be used to express eukaryotic DNA in a variety of hosts such as bacteria, bluegreen algae, plant cells, insect cells and animal cells. Specifically designed vectors allow the shuttling of DNA between hosts such as bacteria-yeast or bacteria-animal cells. An appropriately constructed expression vector should contain: an origin of replication for autonomous replication in host cells, selectable markers, a limited number of useful restriction enzyme sites, a potential for high copy number, and active promoters. A promoter is defined as a DNA sequence that directs RNA polymerase to bind to DNA and initiate RNA synthesis. A strong promoter is one which causes mRNAs to be initiated at high frequency. Expression vectors may include, but are not limited to, cloning vectors, modified cloning vectors, specifically designed plasmids or viruses.

Commercially available mammalian expression vectors which may be suitable for recombinant UCP3 expression, include but are not limited to, pcDNA3.1 (Invitrogen), pBlueBacHis2 (Invitrogen), pLITMUS28, pLITMUS29, pLITMUS38 and pLITMUS39 (New England Biolabs), pcDNA1, pcDNA1amp (Invitrogen), pcDNA3 (Invitrogen), pMC1neo (Stratagene), pXT1 (Stratagene), pSG5 (Stratagene), EBO-pSV2-neo (ATCC 37593) pBPV-1(8-2) (ATCC 37110), pdBPV-MMTneo(342-12) (ATCC 37224), pRSVgpt (ATCC 37199), pRSVneo (ATCC 37198), pSV2-dhfr (ATCC 37146), pUCTag (ATCC 37460), and lZD35 (ATCC 37565).

A variety of bacterial expression vectors may be used to express recombinant UCP3 in bacterial cells. Commercially available

bacterial expression vectors which may be suitable for recombinant UCP3 expression include, but are not limited to pCR2.1 (Invitrogen), pET11a (Novagen), lambda gt11 (Invitrogen), pcDNAII (Invitrogen), pKK223-3 (Pharmacia).

5 A variety of fungal cell expression vectors may be used to express recombinant UCP3 in fungal cells. Commercially available fungal cell expression vectors which may be suitable for recombinant UCP3 expression include but are not limited to pYES2 (Invitrogen), *Pichia* expression vector (Invitrogen).

10 A variety of insect cell expression vectors may be used to express recombinant receptor in insect cells. Commercially available insect cell expression vectors which may be suitable for recombinant expression of UCP3 include but are not limited to pBlueBacIII and pBlueBacHis2 (Invitrogen).

15 An expression vector containing DNA encoding a UCP3-like protein may be used for expression of UCP3 in a recombinant host cell. Recombinant host cells may be prokaryotic or eukaryotic, including but not limited to bacteria such as *E. coli*, fungal cells such as yeast, mammalian cells including but not limited to cell lines of human, 20 bovine, porcine, monkey and rodent origin, and insect cells including but not limited to *Drosophila* and silkworm derived cell lines. Cell lines derived from mammalian species which may be suitable and which are commercially available, include but are not limited to, L cells L-M(TK-) (ATCC CCL 1.3), L cells L-M (ATCC CCL 1.2), Saos-2 (ATCC HTB-85), 293 (ATCC CRL 1573), Raji (ATCC CCL 86), CV-1 (ATCC CCL 70), COS-1 (ATCC CRL 1650), COS-7 (ATCC CRL 1651), CHO-K1 (ATCC CCL 61), 3T3 (ATCC CCL 92), NIH/3T3 (ATCC CRL 1658), HeLa (ATCC CCL 2), C127I (ATCC CRL 1616), BS-C-1 (ATCC CCL 26) and MRC-5 (ATCC CCL 171).

30 The expression vector may be introduced into host cells via any one of a number of techniques including but not limited to transformation, transfection, protoplast fusion, and electroporation. The expression vector-containing cells are individually analyzed to determine whether they produce UCP3 protein. Identification of UCP3 35 expressing cells may be done by several means, including but not

limited to immunological reactivity with anti-UCP3 antibodies, and the presence of host cell-associated UCP3 activity.

The cloned UCP3 cDNA obtained through the methods described above may be recombinantly expressed by molecular cloning into an expression vector (such as pcDNA3.1, pCR2.1, pBlueBacHis2 and pLITMUS28) containing a suitable promoter and other appropriate transcription regulatory elements, and transferred into prokaryotic or eukaryotic host cells to produce recombinant UCP3. Techniques for such manipulations can be found described in Sambrook, et al., *supra*, are discussed at length in the Example section and are well known and easily available to the artisan of ordinary skill in the art.

Expression of UCP3 DNA may also be performed using *in vitro* produced synthetic mRNA. Synthetic mRNA can be efficiently translated in various cell-free systems, including but not limited to wheat germ extracts and reticulocyte extracts, as well as efficiently translated in cell based systems, including but not limited to microinjection into frog oocytes, with microinjection into frog oocytes being preferred.

To determine the UCP3 cDNA sequence(s) that yields optimal levels of UCP3 protein, UCP3 cDNA molecules including but not limited to the following can be constructed: the full-length open reading frame of the UCP3 cDNA and various constructs containing portions of the cDNA encoding only specific domains of the protein or rearranged domains of the protein. All constructs can be designed to contain none, all or portions of the 5' and/or 3' untranslated region of UCP3. UCP3 activity and levels of protein expression can be determined following the introduction, both singly and in combination, of these constructs into appropriate host cells. Following determination of the UCP3 cDNA cassette yielding optimal expression in transient assays, this UCP3 cDNA construct is transferred to a variety of expression vectors (including recombinant viruses), including but not limited to those for mammalian cells, plant cells, insect cells, oocytes, bacteria, and yeast cells.

Levels of UCP3 protein in host cells is quantified by a variety of techniques including, but not limited to, immunoaffinity and/or

ligand affinity techniques. UCP3-specific affinity beads or UCP3-specific antibodies are used to isolate ³⁵S-methionine labeled or unlabelled UCP3 protein. Labeled UCP3 protein is analyzed by SDS-PAGE. Unlabelled UCP3 protein is detected by Western blotting, ELISA or RIA assays
5 employing UCP3 specific antibodies.

Following expression of UCP3 in a host cell, UCP3 protein may be recovered to provide UCP3 in active form. Several UCP3 purification procedures are available and suitable for use. Recombinant UCP3 may be purified from cell lysates and extracts, or from
10 conditioned culture medium, by various combinations of, or individual application of salt fractionation, ion exchange chromatography, size exclusion chromatography, hydroxylapatite adsorption chromatography and hydrophobic interaction chromatography.

In addition, recombinant UCP3 can be separated from
15 other cellular proteins by use of an immuno-affinity column made with monoclonal or polyclonal antibodies specific for full length UCP3, or polypeptide fragments of UCP3. Additionally, polyclonal or monoclonal antibodies may be raised against a synthetic peptide (usually from about 9 to about 25 amino acids in length) from a portion of the protein as
20 disclosed in SEQ ID NO:12. Monospecific antibodies to UCP3 are purified from mammalian antisera containing antibodies reactive against UCP3 or are prepared as monoclonal antibodies reactive with UCP3 using the technique of Kohler and Milstein (1975, *Nature* 256: 495-497). Monospecific antibody as used herein is defined as a single
25 antibody species or multiple antibody species with homogenous binding characteristics for UCP3. Homogenous binding as used herein refers to the ability of the antibody species to bind to a specific antigen or epitope, such as those associated with the UCP3, as described above. UCP3 specific antibodies are raised by immunizing animals such as mice,
30 rats, guinea pigs, rabbits, goats, horses and the like, with an appropriate concentration of UCP3 or UCP3 synthetic peptide either with or without an immune adjuvant.

Preimmune serum is collected prior to the first immunization. Each animal receives between about 0.1 mg and about
35 1000 mg of UCP3 associated with an acceptable immune adjuvant. Such

acceptable adjuvants include, but are not limited to, Freund's complete, Freund's incomplete, alum-precipitate, water in oil emulsion containing *Corynebacterium parvum* and tRNA. The initial immunization consists of the UCP3 protein or UCP3 synthetic peptide in, preferably, Freund's complete adjuvant at multiple sites either subcutaneously (SC), intraperitoneally (IP) or both. Each animal is bled at regular intervals, preferably weekly, to determine antibody titer. The animals may or may not receive booster injections following the initial immunization. Those animals receiving booster injections are generally given an equal amount of UCP3 in Freund's incomplete adjuvant by the same route. Booster injections are given at about three week intervals until maximal titers are obtained. At about 7 days after each booster immunization or about weekly after a single immunization, the animals are bled, the serum collected, and aliquots are stored at about -20°C.

Monoclonal antibodies (mAb) reactive with UCP3 are prepared by immunizing inbred mice, preferably Balb/c, with UCP3. The mice are immunized by the IP or SC route with about 1 mg to about 100 mg, preferably about 10 mg, of UCP3 in about 0.5 ml buffer or saline incorporated in an equal volume of an acceptable adjuvant, as discussed above. Freund's complete adjuvant is preferred. The mice receive an initial immunization on day 0 and are rested for about 3 to about 30 weeks. Immunized mice are given one or more booster immunizations of about 1 to about 100 mg of UCP3 in a buffer solution such as phosphate buffered saline by the intravenous (IV) route. Lymphocytes, from antibody positive mice, preferably splenic lymphocytes, are obtained by removing spleens from immunized mice by standard procedures known in the art. Hybridoma cells are produced by mixing the splenic lymphocytes with an appropriate fusion partner, preferably myeloma cells, under conditions which will allow the formation of stable hybridomas. Fusion partners may include, but are not limited to: mouse myelomas P3/NS1/Ag 4-1; MPC-11; S-194 and Sp 2/0, with Sp 2/0 being preferred. The antibody producing cells and myeloma cells are fused in polyethylene glycol, about 1000 mol. wt., at concentrations from about 30% to about 50%. Fused hybridoma cells are selected by growth in

hypoxanthine, thymidine and aminopterin supplemented Dulbecco's Modified Eagles Medium (DMEM) by procedures known in the art. Supernatant fluids are collected from growth positive wells on about days 14, 18, and 21 and are screened for antibody production by an
5 immunoassay such as solid phase immunoradioassay (SPIRA) using UCP3 as the antigen. The culture fluids are also tested in the Ouchterlony precipitation assay to determine the isotype of the mAb. Hybridoma cells from antibody positive wells are cloned by a technique such as the soft agar technique of MacPherson, 1973, Soft Agar
10 Techniques, in *Tissue Culture Methods and Applications*, Kruse and Paterson, Eds., Academic Press.

Monoclonal antibodies are produced *in vivo* by injection of pristine primed Balb/c mice, approximately 0.5 ml per mouse, with about 2×10^6 to about 6×10^6 hybridoma cells about 4 days after priming.
15 Ascites fluid is collected at approximately 8-12 days after cell transfer and the monoclonal antibodies are purified by techniques known in the art.

In vitro production of anti-UCP3 mAb is carried out by growing the hybridoma in DMEM containing about 2% fetal calf serum to obtain sufficient quantities of the specific mAb. The mAb are purified
20 by techniques known in the art.

Antibody titers of ascites or hybridoma culture fluids are determined by various serological or immunological assays which include, but are not limited to, precipitation, passive agglutination,
25 enzyme-linked immunosorbent antibody (ELISA) technique and radioimmunoassay (RIA) techniques. Similar assays are used to detect the presence of UCP3 in body fluids or tissue and cell extracts.

It is readily apparent to those skilled in the art that the above described methods for producing monospecific antibodies may be
30 utilized to produce antibodies specific for UCP3 polypeptide fragments, or full-length UCP3 polypeptide.

UCP3 antibody affinity columns are made by adding the antibodies to Affigel-10 (Biorad), a gel support which is pre-activated with N-hydroxysuccinimide esters such that the antibodies form
35 covalent linkages with the agarose gel bead support. The antibodies are

then coupled to the gel via amide bonds with the spacer arm. The remaining activated esters are then quenched with 1M ethanolamine HCl (pH 8). The column is washed with water followed by 0.23 M glycine HCl (pH 2.6) to remove any non-conjugated antibody or
5 extraneous protein. The column is then equilibrated in phosphate buffered saline (pH 7.3) and the cell culture supernatants or cell extracts containing UCP3 or UCP3 fragments are slowly passed through the column. The column is then washed with phosphate buffered saline until the optical density (A_{280}) falls to background, then the protein is
10 eluted with 0.23 M glycine-HCl (pH 2.6). The purified UCP3 protein is then dialyzed against phosphate buffered saline.

The novel UCP3 of the present invention is suitable for use in an assay procedure for the identification of compounds which modulate UCP3 activity. Modulating UCP3 activity, as described herein
15 includes the inhibition or activation of the protein and also includes directly or indirectly affecting the cell cycle regulatory properties associated with UCP3 activity. Compounds which modulate UCP3 activity include agonists, antagonists, inhibitors, activators, and compounds which directly or indirectly affect regulation of the UCP3
20 activity and/or the UCP3 uncoupling activity.

The UCP3 of the present invention may be obtained from both native and recombinant sources for use in an assay procedure to identify UCP3 modulators. In general, an assay procedure to identify UCP3 modulators will contain the UCP3-protein of the present
25 invention, and a test compound or sample which contains a putative UCP3 modulator. The test compounds or samples may be tested directly on, for example, purified UCP3 protein whether native or recombinant, subcellular fractions of UCP3-producing cells whether native or recombinant, and/or whole cells expressing the UCP3 whether native or
30 recombinant. The test compound or sample may be added to the UCP3 in the presence or absence of a known UCP3 modulator. The modulating activity of the test compound or sample may be determined by, for example, analyzing the ability of the test compound or sample to bind to UCP3 protein, activate the protein, inhibit UCP3 activity, inhibit

Therefore, the present invention also relates to subcellular membrane fractions of the recombinant host cells (both prokaryotic and eukaryotic as well as both stably and transiently transformed cells) comprising the nucleic acids of the present invention. These subcellular membrane fractions will comprise UCP3 at levels substantially above wild type levels and hence will be useful in various assays described throughout this specification.

The identification of modulators of UCP3 activity will be useful in treating disease states involving, as an example but not a limitation, obesity and diabetes, by manipulating the interrelated process of balancing food intake, energy expenditure and glucose metabolism within the patient. Therefore, modulators to treat hyperactive conditions of energy expenditure which originate in the mitochondria of skeletal muscle will also be within the purview of the present invention.

The present invention is also directed to methods for screening for compounds which modulate the expression of DNA or RNA encoding a UCP3 protein of the present invention or which modulates the function of a such a UCP3 protein. Compounds which modulate these activities may be DNA, RNA, peptides, proteins, or non-proteinaceous organic molecules. Compounds may modulate by increasing or attenuating the expression of DNA or RNA encoding the UCP3 protein, or the function of a UCP3 protein. Compounds that modulate the expression of DNA or RNA encoding the UCP3 protein or the biological function thereof may be detected by a variety of assays. The assay may be a simple "yes/no" assay to determine whether there is a change in expression or function. The assay may be made quantitative by comparing the expression or function of a test sample with the levels of expression or function in a standard sample. Kits containing modified UCP3, antibodies to UCP3, or modified UCP3 protein may be prepared by known methods for such uses.

The DNA molecules, RNA molecules, recombinant protein and antibodies of the present invention may be used to screen and

30

measure levels of UCP3 DNA, RNA or protein. The recombinant proteins, DNA molecules, RNA molecules and antibodies lend themselves to the formulation of kits suitable for the detection and typing of UCP3. Such a kit would comprise a compartmentalized carrier
5 suitable to hold in close confinement at least one container. The carrier would further comprise reagents such as recombinant UCP3 protein or anti-UCP3 antibodies suitable for detecting UCP3. The carrier may also contain a means for detection such as labeled antigen or enzyme substrates or the like.

10 Pharmaceutically useful compositions comprising modulators of UCP3 may be formulated according to known methods such as by the admixture of a pharmaceutically acceptable carrier. Examples of such carriers and methods of formulation may be found in Remington's Pharmaceutical Sciences. To form a pharmaceutically
15 acceptable composition suitable for effective administration, such compositions will contain an effective amount of the protein, DNA, RNA, or modified UCP3.

Therapeutic or diagnostic compositions of the invention are administered to an individual in amounts sufficient to treat or diagnose
20 disorders. The effective amount may vary according to a variety of factors such as the individual's condition, weight, sex and age. Other factors include the mode of administration.

The pharmaceutical compositions may be provided to the individual by a variety of routes such as subcutaneous, topical, oral and
25 intramuscular.

The term "chemical derivative" describes a molecule that contains additional chemical moieties which are not normally a part of the base molecule. Such moieties may improve the solubility, half-life, absorption, etc. of the base molecule. Alternatively the moieties may
30 attenuate undesirable side effects of the base molecule or decrease the toxicity of the base molecule. Examples of such moieties are described in a variety of texts, such as Remington's Pharmaceutical Sciences.

Compounds identified according to the methods disclosed herein may be used alone at appropriate dosages. Alternatively, co-

199591
administration or sequential administration of other agents may be desirable.

5 The present invention also has the objective of providing suitable topical, oral, systemic and parenteral pharmaceutical formulations for use in the novel methods of treatment of the present invention. The compositions containing compounds identified according to this invention as the active ingredient can be administered in a wide variety of therapeutic dosage forms in conventional vehicles for administration. For example, the compounds can be administered
10 in such oral dosage forms as tablets, capsules (each including timed release and sustained release formulations), pills, powders, granules, elixirs, tinctures, solutions, suspensions, syrups and emulsions, or by injection. Likewise, they may also be administered in intravenous (both bolus and infusion), intraperitoneal, subcutaneous, topical with or
15 without occlusion, or intramuscular form, all using forms well known to those of ordinary skill in the pharmaceutical arts.

Advantageously, compounds of the present invention may be administered in a single daily dose, or the total daily dosage may be administered in divided doses of two, three or four times daily.
20 Furthermore, compounds for the present invention can be administered in intranasal form via topical use of suitable intranasal vehicles, or via transdermal routes, using those forms of transdermal skin patches well known to those of ordinary skill in that art. To be administered in the form of a transdermal delivery system, the dosage administration will,
25 of course, be continuous rather than intermittent throughout the dosage regimen.

For combination treatment with more than one active agent, where the active agents are in separate dosage formulations, the active agents can be administered concurrently, or they each can be
30 administered at separately staggered times.

The dosage regimen utilizing the compounds of the present invention is selected in accordance with a variety of factors including type, species, age, weight, sex and medical condition of the patient; the severity of the condition to be treated; the route of administration; the
35 renal and hepatic function of the patient; and the particular compound

5

10

15

20

Identification of Partial DNA Fragments Encoding Human UCP3

25

T,0340

GGTGACCTACGACATCCTCAAGGAGAAGCTGCTGGACTACCACCTGCTCACTGACAACCT
 TCCCCTGCCACTTTGTCTCTGCCTTTGGAGCCGGCTTCTGTGCCACAGTGGTGGCATCC
 CCGGTGGACGTGGTGAAGACCCGGTATATGAACTCACCTCCAGGCCAGTACTTCAGCCC
 CCTCGACTGTATGATAAAGATGGTGGCCCAGGAGCGCCACCAGCCTTCTACAAGGGATT
 5 TACACCCTCCTTTTTGCGTTTGGGATCCTGGAACGTGGTGTATGTTTCGTAACCTATGAGC
 AGCTGAAACGGGCCCTGATGAAAGTCCAGATGTTACGGGAATCACCGTTTTGAACAAGA
 CAAGAAGGCCACTGGTAGCTAACGTGTCCGAAACCAGTTAAGAATGGAAG
 (SEQ ID NO:1).

10 An "N" residue is added at base number 249 in order to preserve the open
 reading frame of this EST. This corrected version of AA192136 is as
 follows:

T,0341

GGTGACCTACGACATCCTCAAGGAGAAGCTGCTGGACTACCACCTGCTCACTGACAACCT
 TCCCCTGCCACTTTGTCTCTGCCTTTGGAGCCGGCTTCTGTGCCACAGTGGTGGCATCC
 CCGGTGGACGTGGTGAAGACCCGGTATATGAACTCACCTCCAGGCCAGTACTTCAGCCC
 15 CCTCGACTGTATGATAAAGATGGTGGCCCAGGAGCGCCACCAGCCTTCTACAAGGGATT
 TACACCCTCCTTNTTTGCGTTTGGGATCCTGGAACGTGGTGTATGTTTCGTAACCTATGAG
 CAGCTGAAACGGGCCCTGATGAAAGTCCAGATGTTACGGGAATCACCGTTTTGAACAAG
 ACAAGAAGGCCACTGGTAGCTAACGTGTCCGAAACCAGTTAAGAATGGAAG(SEQ ID
 NO:2).

20 SEQ ID NO:1 is a component of Stratagene® muscle cDNA library
 937209 and is identified as cDNA Image Clone No. 628529. This
 Stratagene muscle cDNA library was generated by standard methods
 and cDNAs are cloned unidirectionally into EcoRI/XhoI digested
 pBS(SK-) plasmid. The mRNA for this library was isolated from the
 25 skeletal muscle of an adult human patient suffering from malignant
 hyperthermia. This cDNA clone is publically available by Genbank
 Accession No. AA192136, Image Clone ID No. 628529, and Washington
 University Clone ID No. zq02d09.r1. This construct is available from
 Research Genetics, Inc., 2130 Memorial Parkway SW, Huntsville, AL
 30 35801 (<http://www.resgen.com>).

SEQ ID NO:1 was used as a query to search Genbank with Blastn.
 A second EST, with accession number Z28895 (and assigned Genbank
 number HSB6C051) was identified and is disclosed throughout this
 specification as SEQ ID NO:3, which is:

T₁0350
CGAGCAGCTGAAACGGGCCCTGATGAAAGTCCAGATGTTACGGGNATCACCGTTTTGAA
CAAGACAAGAAGGCCACTGGTAGCTAACGTNTCCGAAACCAGTTAAGANTGGAAGAAAA
CGGTCCATCCACGNACACATGGACACAGACCCACACATNTT (SEQ ID NO:3).

The EST disclosed as SEQ ID NO:2 shows an open reading frame
5 encoding a novel protein fragment showing homology to human UCP1
and human UCP2. SEQ ID NO:2 represents the 5' portion of the cDNA
Image Clone No. 628529. The 3' sequence read of cDNA Image Clone
No. 628529 has the accession number aa192553, which is as follows:

TTTTTTTTGTCTTACTCCACACCTAAGGTGGAANTTCTTTTATTGAGTCATAATAAT
10 TTCCCGAGAATTCCGAGTCCTGCTACTTTAGGTTCTTGCCCAGGAATCCACCTCTTTTC
CCCCAAGCCCCAACAAATCCTTTGAGGTACTCATGATTGAGCGCGTGGTGGGGGGGGGTGG
GGAAGAGGCTGCATGGGGGTGGGGCTCCTGTGGCTTCACGTCATCCACTGTCACCTCTG
GTCCCCAAGTCTCTGGATCCTTTGGTCTCACCTCTAGACAACCGCGGGGTTCAAACCT
TCTTCCCTGGCAACTCCTCTCTGTCCCGACAAAATCTCTCCCAAGGCATTGTCTTGTGA
15 GTTAGATTACACAGAGCTTTTGCTTTTATAAAGTGCGTTCATGCCAGCTTCTCACTT
GCATGTCATAGCACCCCTGGTGAGGTGGACAGGGAAGGGATGGCTCCCTCCATTTTGTGA
GGAAAGTNGGGG (SEQ ID NO:4).

20

EXAMPLE 2:

Isolation and Characterization of DNA Fragments Encoding Full-Length Human UCP3

25 The exemplified full length cDNA of the present invention was
generated by a novel method of cDNA screening as described in the
Detailed Description of the Invention.

30 The foregoing process is exemplified for isolation and
characterization of a full-length cDNA encoding human UCP3. Briefly,
a random oligo dT primed fetal brain cDNA library consisting of
approximately 4.0 million primary clones was constructed in the
plasmid vector pBluescript (Stratagene, LaJolla, CA). The primary
clones were subdivided into 188 pools with each pool containing
approximately 20,000 clones. Each pool was amplified separately and
the resulting plasmid pools were collected and transferred into two
35 96-well plates.

25

Construction of a Yeast Expression Vector and Transformed Yeast Cell Line Expressing Human UPC3

30

5'CATTCCTCGAGCTACCAGTGGCCTTCTTG3', SEQ ID NO:14) were used to generate a PCR product of human UCP3 tagged with EcoRI and XhoI sites at the 5' and 3' ends, respectively, using pCR2.1-UCP3.1 as the template. The PCR product was digested with EcoRI and XhoI and purified from an agarose gel. This fragment was ligated into a yeast expression vector pYES2 (Invitrogen, San Diego, CA) that was also digested with EcoRI and XhoI such that UCP3 is under the control of *S. cerevisiae* GAL1 promoter. Multiple recombinants isolated after transformation into *E. coli* were picked and sequenced using three UCP3 internal primers in combination with two pYES2 vector primers (5'CCCGGATCGGACTACTAGCA, (SEQ ID NO:15); 5'GGGGGGGAGGGCGTGAATGTAA; [SEQ ID NO:16]). Mutation-free clones of pYES2-UCP3 were then transformed into two *Saccharomyces cerevisiae* strains INVSc1 (*MATa*, *his3D1*, *leu2*, *trp1-289*, *ura3-52*) and YPH499(*MATa*, *ura3-52*, *lys2-801*, *ade2-101*, *trp1-D63*, *his3-D200*, *leu2- D1*) using lithium acetate method and selected on Sc-ura media. The expression of UCP3 was achieved by induction in Sc-ura media with 2% galactose and 3% glycerol.

The diploid *Saccharomyces cerevisiae* strain W303 (*MATa/a*, *can1-100*, *ade2-10*, *his3-11,-15*, *leu2-2,-112*, *trp1-D1*, *ura3-1*) was also used for the expression of UCP3. Human UCP3 was expressed under the control of GAL1 promoter in pYES2 (Invitrogen, San Diego, CA). Constructs were made directly in yeast using a gap repair strategy. The entire open reading frame of human UCP3 was amplified using a pair of primers that incorporated a yeast translation start site (ATAATG). Taq Additive (Stratagene, La Jolla, CA) was included in PCR reaction to enhance amplification fidelity. The PCR product was fused to GAL1 at the 5' end and to CYC1 terminator at the 3' end by co-transforming the PCR product and pYES2 followed by selection on Sc-Ura medium. More than 50% of the transformants contained the expected GAL1-UCP3 plasmids. Consistent results were obtained from multiple independent clones for growth and flow cytometry analysis. For the growth assay, the W303 strain containing the vector alone or UCP3 expression plasmid was streaked onto Sc-Ura media with 3%

glycerol and 2% galactose and incubated at 30°C for 48 hours. To measure mitochondria membrane potential, cells were grown in Sc-Ura media supplemented with raffinose. Galactose was added to induce the expression of UCP3 for five hours. Fluorescent probe DiOC6 (3,3'-
5 dihexyloxacarbocyanine iodine) was added after cell concentration was adjusted to two millions per ml. Data were collected and analyzed as described (Bouillaud et al., 1994, *EMBO J.* 13: 1990-1997). A total 10,000 objects were taken for the histograms. For the display of fluorescence intensity, a three-order logarithmic scale was used.

10 The biochemical activity of UCP1 and UCP2 have previously been studied by measuring the defects of aerobic growth in yeast expressing UCPs. To test that UCP3 encodes an uncoupling protein for mitochondria, UCP3 was expressed under the control of a tightly inducible promoter *GAL1* in the yeast system as previously described for
15 UCP1 and UCP2 (Fleury et al., 1997, *Nature Genetics* 15: 269-272; Gimeno et al., 1997, *Diabetes* 46, 900-906). Induction with galactose resulted in a significant decrease in the growth rate of strains containing UCP3 relative to cells with vector alone. To confirm more directly that UCP3 reduced the mitochondria membrane potential, the
20 fluorescent intensity of cells labeled with the potential sensitive dye DiOC6 was measured using flow cytometry analysis. Induction of UCP3 expression by galactose significantly reduced the membrane potential of mitochondria compared with strains containing vector alone (Figure 8A and 8C). In addition, no membrane potential shift was observed when
25 UCP3 expression was not induced (Figure 8B). Two membrane potential peaks were observed after UCP3 induction (Figure 8C). Similar results were observed for UCP2 (Fleury et al., 1997, *Nature Genetics* 15: 269-272). The most likely explanation for this observation is that one peak represents cells with reduced membrane potential due to expression of
30 UCP3 while the other peak represents cells with normal membrane potential due to the loss of UCP3 expression. When UCP2 was cloned into the same vector as UCP3 and transformed into the same yeast strain, UCP3 reduced membrane potential to a greater extent compared with UCP2. The data indicate that UCP3 encodes an uncoupling protein
35 that is able to decrease the membrane potential of mitochondria.

EXAMPLE 4:

Tissue Distribution Of UCP3 Expression

5 Human multiple tissue Northern Blot #7760-1, Human
Brain Northern Blot II #7755-1, Human Brain Northern Blot III
#7750-1, and Human multiple tissue Northern Dot Blot were purchased
from Clontech. The probe was made by PCR amplification of coding
region of UCP3 (SEQ ID NO:11). This fragment was labeled with ³²P and
10 used to probe the Northern blot as described in the Clontech procedures,
which are well known in the art. The blots were exposed to X-ray film
with an intensifying screen at -70 °C for times as indicated in the figure.
Figure 4A-4C is an autoradiograph which shows that UCP3 is highly
expressed in skeletal muscle as a transcript of ~2.4 kb (Figure 4A & B).
15 Two minor transcripts, ~6.0 kb and 7.5 kb are also expressed in skeletal
muscle (Figure 4A). In addition, UCP3 is also weakly expressed as a
transcript of ~2.4 kb in the heart. Probing with human beta-actin probe
(Figure 4C) shows that all the lanes have approximately equal amount of
total polyA RNA, indicating that detection of UCP3 only in skeletal
20 muscle and in the heart is not due to the lack of RNA in the other tissue
lanes. Also, In the brain, ucp3 is most abundantly expressed in the
corpus collosum as a transcript of ~2.4 kb.

The expression pattern of UCP3 appears to be unique, because it is
abundantly and nearly exclusively expressed in skeletal muscle. Even
25 though UCP3 expression was also detected in the heart and brain, it was
found at much lower levels. Skeletal muscle is responsible for more
than half of energy expenditure at resting stage. Differences in energy
expenditure of skeletal muscle can explain most of the variation in basal
metabolic rate between individuals (Ravussin and Bogardus, 1992, *Am J*
30 *Clin Nutr* 55: 242S-245S). The proton leak was shown to account for
around one-half of the resting respiration rate of perfused rat skeletal
muscle (Rolfe and Brand, 1996, *Am J Physiol* 271: C1380-1389).
Furthermore, free fatty acids-activated uncoupling in skeletal muscle

5

Characterization and Chromosomal Localization of Genes Encoding UCP3

15

20

25

30

chromosomal region in both human and mouse, while human UCP1 was mapped to 4q31. The chromosomal regions where UCP2 resides are linked to obesity and hyperinsulinemia in both human and mouse, and so UCP2 was hypothesized to be a candidate gene for these diseases (Taylor and Phillips, 1996, *Genomics* 34, 389-398; Fleury et al., 1997, *Nature Genetics* 15: 269-272). The close location of UCP3 to UCP2 suggests that UCP3 is also a candidate gene for obesity and hyperinsulinemia. In addition, the close homology and location between UCP2 and UCP3 suggested that the two genes arose from a gene duplication event after UCP1 was separated.

Alignment of the six protein sequences of UCP1 to UCP3 of human and mouse reveals a highly clustered pattern of the conserved residues. The transmembrane domains and their immediate following sequences are much more conserved than the sequences between the transmembrane domains. Moreover, each allbnb motif following the first transmembrane domain of each repeat unit is perfectly conserved among the six proteins. Thus, UCP3 is a typical member of the mitochondria transport protein superfamily. UCP1 activity is inhibited by nucleotides but activated by free fatty acids. Affinity labeling and site-directed mutagenesis demonstrated that the region between transmembrane domain 5 and 6 bound to nucleotides (Mayinger and Klingenberg, 1992, *Biochemistry* 31: 10536-10543; Winkler and Klingenberg, 1992, *Eur J Biochem* 203: 295-304). Particularly, deletion of the putative nucleotide binding motif (amino acid residue 261 to 269) renders UCP1 resistant to nucleotide-mediated inhibition (Bouillaud et al., 1994, *EMBO J* 13, 1990-1997). Substitution of Phe 267 by a Tyr also results in higher uncoupling activity (Bouillaud et al., id.). Both UCP2 and UCP3 contain a Tyr instead of a Phe at this position, suggesting that UCP2 and UCP3 has higher basal activity. On the other hand, UCP1 and UCP3 both have a Thr at the position corresponding residue 264 of UCP1 whereas UCP2 has an Arg. The presence of a positively charged residue in this motif may imply a higher affinity of UCP2 for nucleotides and, consequently, increased sensitivity to nucleotide-mediated inhibition. The very C-terminus of UCP1 was shown to be important for activation by fatty acids (Gonzalez-Barroso et al., 1996, *Eur J Biochem*

15

Adenovirus-Mediated Leptin Expression

20

25

30

It has been reported that adenovirus-mediated leptin expression leads to significant reduction of body weight in ob/ob mice. Given the abundance of UCP3 in skeletal muscle, the effect of leptin treatment on UCP3 expression in skeletal muscle was tested by injecting

5 adenoviruses expressing leptin into ob/ob mice and examining UCP3 expression after one week. Injection of adenoviruses expressing leptin, either helper-dependent (HD-leptin) or first generation (Ad-leptin), led to significant weight loss (approximately 15%) after one week while

10 injection of adenoviral vector expressing b-galactosidase (Adb-gal) had no significant effect on body weight (Figure 9A). Measurement of serum leptin levels at the end of one week confirmed high levels of leptin production in mice injected with HD-leptin (26.7 ± 10 ng/ml) or Ad-leptin (21.8 ± 16.7 ng/ml) but not in mice injected with Adb-gal (1.56 ± 0.05 ng/ml) or buffer alone (2 ± 0.17 ng/ml). Wild type lean mice have leptin

15 levels at approximately 4.5 ng/ml. UCP3 levels were then checked in skeletal muscle of these animals. Mice treated with adenovectors expressing leptin, either Ad-leptin or HD-leptin, showed approximately an increase of 80% and 70%, respectively, in UCP3 RNA level (Figure 9B, $p=0.2$ vs. buffer-injected mice). In contrast, mice treated with Adb-gal

20 did not show significant change in UCP3 expression. If the average UCP3 level of animals showing weight loss (leptin-treated, $n=6$) was compared with that of animals showing no weight loss (b-galactosidase treated or untreated, $n=4$), the increase in UCP3 expression after leptin treatment was statistically significant ($p=0.01$). Thus, leptin treatment

25 leads to increased UCP3 expression in skeletal muscle in ob/ob mice.

Leptin plays a pivotal role in the regulation of food intake and energy expenditure. Leptin-deficient ob/ob mice are hyperphagic and hypometabolic, and leptin treatment led to decreased food intake and increased metabolic rate (Halaas et al., 1995, *Science* 269: 543-546, Hwa et al., 1997, *Am J Physiol* 272, R1204-1209; Hwa et al., 1996 *Horm Metab Res* :28: 659-663). These results suggest that UCP3 may be involved in the increase of metabolic rate post-leptin treatment. Leptin treatment led to increased expression of UCP2 in pancreatic islets and adipose tissue (Zhou et al., 1997). Thus, both UCP2 and UCP3 are likely to be

35 involved in the increase of metabolic rate post-leptin treatment.

EXAMPLE 7:

Isolation and Characterization of DNA Fragments Encoding Full-Length Mouse UCP3

5

Mouse UCP3 was isolated by semi-nested PCR followed by 5' and 3' race. Three degenerate PCR primers were designed based on the conserved regions of the three uncoupling proteins: F441, 5'-CCNCTGGAYACNGCYAA-3' (SEQ ID NO:19); F755, 5'-CAGCCCCACNGANGTNGT-3' (SEQ ID NO:20); and R1055, 5'-TTCACCACRTCACNACNGG-3' (SEQ ID NO:21). First round PCR was carried out on mouse skeletal muscle cDNA with the primer pair F441+R1055. Second round PCR was carried out with the primer pair F755+R1055. All PCR reactions were carried out using AmpliTaq Gold (Perkin Elmer, NJ). The degerate PCR product was cloned into the vector pCR2.1 (Invitrogen, CA) and sequenced. Analysis of the sequences identified a partial cDNA sequence that was highly homologous to human UCP3. This partial sequence were then used to isolate the full-length sequence of mouse UCP3 by 5' and 3' race using a Marathon race kit from Clontech (Clontech Laboratories, CA). DNA sequences were all determined using the ABI Prism dye terminator system on ABI 377 machines. The full length cDNA encoding mouse UCP3 (SEQ ID NO:17) is shown in Figure 6A and 6B and the amino acid sequence of mouse UCP3 (SEQ ID NO:18) is shown in Figure 7.

25